Oligonucleotide primers used and PCR cycling conditions for de amplification and sequencing of six housekeeping genes using in the MLST of *Yersinia ruckeri*

Gene	Primer (5'-3')	Length (bp)	PCR cycling	Reference
*gln*A	Forward: CGATTGGTGGCTGGAAAGGC Reverse: TTGGTCATRGTRTTGAAGCG	530	5 min 94 ºC, 35 x (45 s 94 ºC, 45 s 51 ºC, 1 min 72 ºC), 5 min 72 ºC	Kotetishvili et al., 2005
*gyr*B	Forward: CGGCGGTTTGCAYGGYGTRGG Reverse: CAGSGTRCGRGTCATYGCCG	545	5 min 94 ºC, 35 x (45 s 94 ºC, 45 s 62 ºC, 1 min 72 ºC), 5 min 72 ºC	Kotetishvili et al., 2005
*rec*A	Forward: GGGCCAAATTGAAAARCARTTCGG Reverse: CGCCRATYTTCATRCGRATYTGGT	560	5 min 94 ºC, 35 x (45 s 94 ºC, 45 s 51 ºC, 1 min 72 ºC), 5 min 72 ºC	Kotetishvili et al., 2005
*dna*J	Forward: ATGGCGAAGAGAGACTATTAC Reverse: AAGCTTTTAGAGCGTGGGTGT	1024	5 min 94 ºC, 35 x (45 s 94 ºC, 45 s 51 ºC, 1 min 72 ºC), 5 min 72 ºC	Hurst et al., 2010
*thr*A	Forward: TGACCATCGCCGTTAT Reverse: GCTTTTGTTGGCGTAC	693	5 min 94 ºC, 35 x (45 s 94 ºC, 45 s 51 ºC, 1 min 72 ºC), 5 min 72 ºC	Tinsley et al., 2009
Y-HSP60	Forward: GACGTNGTAGAAGGTATGYAG Reverse: CGCCGCCAGCCAGTTTAGC	565	5 min 94 ºC, 35 x (45 s 94 ºC, 45 s 51 ºC, 1 min 72 ºC), 5 min 72 ºC	Kotetishvili et al., 2005

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